

CCR Research Orientation Checklist

Name:

ACTIVITY	DATE COMPLETED
General	
NCI Orientation	
CITI Biomedical 101	
CITI GCP	
Safe Shipping and Handling of Infectious and Biological	
Clinical Research Modules	
Clinical Trial Design	
Protocol Development	
Phase 0 Clinical Trials	
Informed Consent Process	
Roles and Responsibilities of the Research Team	
Roles and Responsibilities of the Sponsor	
Clinical Data Management	
Documentation in Clinical Research	
Documenting, Recording, and Reporting of Adverse Events and Unanticipated Problems	
Regulatory Binder	
RECIST: Applying the Rules	
Monitoring & Auditing of Clinical Trials	
Independent Readings	
CCR Policies and SOPs	

ACTIVITY	DATE SCHEDULED	DATE COMPLETED
Orientation Classes		
Protocol Analysis Exercise		
PSO orientation & iRIS training		
C3D training and J-review training		